52044/CAB/R2682

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neurons relative to a control non-transgenic mouse.

WHAT IS CLAIMED IS:

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1. A transgenic mouse whose genome comprises a transgene encoding human small conductance calcium-activated potassium (SK) channel protein, splice variant B1 ("SK3-1B"), wherein the transgene is operably linked to a neuron-specific promoter, and wherein expression of the transgene results in ataxia.

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2. The transgenic mouse of claim 1, wherein the expression of the transgene is limited to neurons.

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3. The transgenic mouse of claim 1, wherein the mouse is fertile and transmits the SK3-1B transgene to its offspring.

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4. The transgenic mouse of claim 1, wherein the SK3-1B transgene has been introduced into an ancestor of said mouse at an embryonic stage.

human SK3-1B transgene.

The transgenic mouse of claim 1, wherein the mouse is hemizygous for the

The transgenic mouse of claim 1, wherein the mouse overexpresses SK3-1B in

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6. The transgenic mouse of claim 1, wherein the mouse is homozygous for the human SK3-1B transgene.

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8. The transgenic mouse of claim 1, wherein the promoter is a Thy1.2-SX promoter.

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52044/CAB/R2682

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- 9. A transgenic mouse whose genome comprises a transgene encoding SK3-1B, wherein the transgene is operably linked to a neuron-specific promoter, and wherein expression of the transgene results in an intention tremor.
- 10. The transgenic mouse of claim 9, wherein the expression of the transgene is limited to neurons.
- 11. The transgenic mouse of claim 9, wherein the mouse is fertile and transmits the SK3-1B transgene to its offspring.
- 12. The transgenic mouse of claim 9, wherein the SK3-1B transgene has been introduced into an ancestor of said mouse at an embryonic stage.
 - 13. The transgenic mouse of claim 9, wherein the mouse is hemizygous for the human SK3-1B transgene.
 - 14. The transgenic mouse of claim 9, wherein the mouse is homozygous for the human SK3-1B transgene.
- 25 The transgenic mouse of claim 9, wherein the mouse overexpresses SK3-1B in neurons relative to a control non-transgenic mouse.
 - 16. The transgenic mouse of claim 9, wherein the promoter is a Thy1.2-SX promoter.
 - 17. A transgenic mouse whose genome comprises a transgene encoding SK3-1B, wherein the transgene is operably linked to a neuron-specific promoter, and wherein expression of the transgene results in hyperexcitable behavior.

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52044/CAB/R2682

18.	The transgenic	mouse	of claim	17,	wherein	the	expression	of the	SK3-1B
transgene is lin	mited to neurons								

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19. The transgenic mouse of claim 17, wherein the mouse is fertile and transmits the SK3-1B transgene to its offspring.

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20. The transgenic mouse of claim 17, wherein the SK3-1B transgene has been introduced into an ancestor of said mouse at an embryonic stage.

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21. The transgenic mouse of claim 17, wherein the mouse is hemizygous for the human SK3-1B transgene.

22. The transgenic mouse of claim 17, wherein the mouse is homozygous for the human SK3-1B transgene.

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23. The transgenic mouse of claim 17, wherein the mouse overexpresses SK3-1B in neurons relative to a control non-transgenic mouse.

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24. The transgenic mouse of claim 17, wherein the promoter is a Thy1.2-SX promoter.

25. A method of screening biologically active agents that facilitate reduction of ataxia in vivo, the method comprising:

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administering a candidate agent to a transgenic mouse according to claim 1, and determining the effect of said agent upon the level of ataxia.

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26. A method of screening biologically active agents that facilitate reduction of intention tremors in vivo, the method comprising:

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administering a candidate agent to a transgenic mouse according to claim 9, and

52044/CAB/R2682

determining the effect of said agent upon the level of intention tremors.

5 27. A method of screening biologically active agents that facilitate improvement in hyperexcitable behavior, the method comprising:

administering a candidate agent to a transgenic mouse according to claim 17, and determining the effect of said agent upon hyperexcitable behavior.